That which is claimed is:

1. A compound having the structure:

$$R^2$$
 R^3
 R^4
 R^5
 $X - OR$

wherein:

A is a C3 up to C8 branched chain alkyl or substituted alkyl group, a C3 up to C7 cycloalkyl or substituted cycloalkyl, an optionally substituted aryl or an optionally substituted heteroaryl,

X is -C(O)- or $-CH_2$ -,

R is methyl or ethyl,

 R^1 is H, hydroxy, alkoxy, benzoyloxy, mesityloxy, or $-OCH_2C(O)OC_2H_5$,

 R^2 is H or R^2 can cooperate with R^3 to form a benzopyran, wherein the pyran ring has the structure:

Me
$$R^6$$
 R^8 R^7

wherein:

R⁶ is not present if the pyran ring is unsaturated, or, if present, is selected from H, -OR, wherein R is alkyl or acyl, or R⁶ can cooperate with R⁷ to form a cyclic acetal, a cyclic ketal, or a cyclopropyl moiety, and

only one of R⁷ and R⁸ is present if the pyran ring is unsaturated, or R⁷ and R⁸ are independently H, carboxyl, cyano, hydroxy, alkoxy, thioalkyl, aryl, or R⁷ and R⁸ taken together comprise a carbonyl oxygen or an oxime nitrogen, or either R⁷ or R⁸ can cooperate with R⁶ to form a cyclic acetal, a cyclic ketal, or a cyclopropyl moiety,

R³ can cooperate with R² to form a benzopyran having the structure set forth above, or R³ is alkenyl, optionally substituted aryl or heteroaryl, or optionally substituted arylalkenyl or heteroarylalkenyl,

R⁵ is H, hydroxy, alkoxy or aryloxy.

- 2. The compound of claim 1 wherein R² and R³ cooperate to form a benzopyran.
- 3. The compound of claim 2 wherein A is cyclopropyl, X is -C(O)-, R^1 is methoxy, R^6 and R^7 are absent, and R^4 , R^5 and R^8 are hydrogen.
- 4. The compound of claim 2 wherein A is cyclopropyl, X is $-CH_2$ -, R^1 is methoxy, R^6 and R^7 are absent, and R^4 , R^5 and R^8 are hydrogen.
- 5. The compound of claim 2 wherein A is cyclohexyl, X is -C(O)-, R^1 is methoxy, R^6 and R^7 are absent, and R^4 , R^5 and R^8 are hydrogen.
- 6. The compound of claim 2 wherein A is phenyl, X is -C(O)-, R^1 is methoxy, R^6 and R^7 are absent, and R^4 , R^5 and R^8 are hydrogen.

- 7. The compound of claim 2 wherein A is phenyl, X is -C(O)-, R¹ is methoxy, R⁶ and R⁷ cooperate to form a dichlorocyclopropyl ring, and R⁴, R⁵ and R⁸ are hydrogen.
- 8. The compound of claim 2 wherein A is cyclohexyl, X is -C(O)-, R¹ is methoxy, R⁶ and R⁷ cooperate to form a dichlorocyclopropyl ring, and R⁴, R⁵ and R⁸ are hydrogen.
 - 9. The compound of claim 1 wherein R^3 is alkenyl.
- 10. The compound of claim 9 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is -CH=CH-C(O)-O-tBu.
 - 11. The compound of claim 1 wherein R³ is optionally substituted aryl or heteroaryl.
- 12. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is phenyl.
- 13. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is p-thiomethyl-phenyl.
- 14. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is m-methoxy-phenyl.
- 15. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is m-acetyl-phenyl.
- 16. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is 5-methyl-2-thiophene-yl.
- 17. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is 5-acetyl-2-thiophene-yl.

- 18. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is 4-dimethylamino-phenyl.
- 19. The compound of claim 11 wherein A is isopropyl, X is -C(O)-, R¹ R², R⁴ and R⁵ are hydrogen, and R³ is 4-dimethylamino-phenyl.
- 20. The compound of claim 11 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is 2,3-(O-CH₂-O)-phenyl.
- 21. The compound of claim 11 wherein A is isopropyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is 2,3-(O-CH₂-O)-phenyl.
- 22. The compound of claim 1 wherein R³ is or optionally substituted arylalkenyl or heteroarylalkenyl.
- 23. The compound of claim 22 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is -CH=CH-phenyl.
- 24. The compound of claim 22 wherein A is isopropyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is -CH=CH-phenyl.
- 25. The compound of claim 22 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is $-\dot{C}H$ =CH-p-methoxy-phenyl.
- 26. The compound of claim 22 wherein A is cyclohexyl, X is -C(O)-, R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is -CH=CH-o-fluoro-phenyl.
- 27. The compound of claim 22 wherein A is isopropyl, X is -C(O)-, R¹ R², R⁴ and R⁵ are hydrogen, and R³ is -CH=CH-o-fluoro-phenyl.

- 28. The compound of claim 22 wherein A is cyclohexyl, X is -C(O), R^1 R^2 , R^4 and R^5 are hydrogen, and R^3 is -CH=CH-m-fluoro-phenyl.
- 29. The compound of claim 22 wherein A is isopropyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is -CH=CH-m-fluoro-phenyl.
- 30. The compound of claim 22 wherein A is cyclohexyl, X is -C(O)-, $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is -CH=CH-p-fluoro-phenyl.
- 31. The compound of claim 22 wherein A is isopropyl, X is -C(O), $R^1 R^2$, R^4 and R^5 are hydrogen, and R^3 is -CH=CH-p-fluoro-phenyl.
- 32. A formulation comprising at least one compound according to claim 1 in a pharmaceutically acceptable carrier therefor.
- 33. A method for modulating process(es) mediated by farnesoid X receptor polypeptides, said method comprising conducting said process(es) in the presence of an effective amount of at least one compound according to claim 1.
- 34. The method of claim 33 wherein said process mediated by farnesoid X receptor is cholesterol metabolism.
- 35. The method of claim 33 wherein said process mediated by farnesoid X receptor is the regulation of lipid homeostasis.
- 36. A method for the treatment of hypercholestemia, said method comprising administering an effective amount of at least one compound according to claim 1 to a subject in need thereof.
- 37. A method for the treatment of cholestasis, said method comprising administering an effective amount of at least one compound according to claim 1 to a subject in need thereof.